

Supplementary information

Title:

Transcriptional repression of *p27* is essential for murine embryonic development

**Youichi Teratake¹, Chisa Ozeki¹, Yuta Hasegawa¹, Yoshiharu Sato², Masayasu Kitahashi²,
Lisa Fujimura³, Haruko Watanabe-Takano³, Akemi Sakamoto^{1,3}, Masafumi Arima²,
Takeshi Tokuhsa², and Masahiko Hatano^{1,3,*}**

**¹Department of Biomedical Science, ²Developmental Genetics, Graduate School of
Medicine, Chiba University,**

³Biomedical Research Center, Chiba University

***To whom correspondence should be addressed: Department of Biomedical Science,
Graduate School of Medicine, Chiba University, 1-8-1 Inohana, Chuo-ku, Chiba city,
Chiba 260-8670, Japan. Tel.: 81-43-226-2950; FAX: 81-43-226-2953; E-mail:
hatanom@faculty.chiba-u.jp**

Running title: Nczf targeted disruption in mice

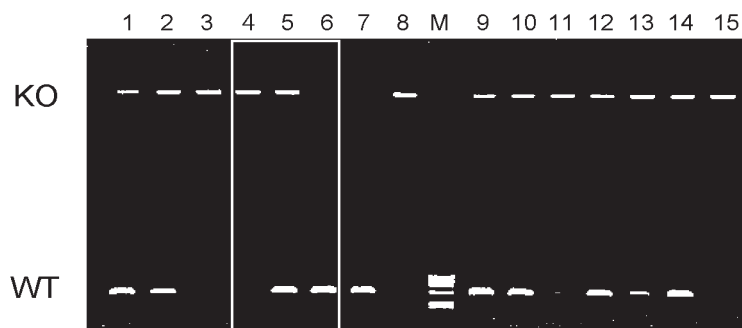


Figure S1

Whole image of gel electrophoresis genotyping Nczf knockout embryos.

Genomic DNA was isolated from E8.5 embryos and PCR was performed with primers specific for wild type and KO allele, respectively. Upper lanes indicate Nczf KO allele (800bp) and lower lanes indicate wild type allele (400bp).

Lanes 4, 5, and 6 were cropped and shown as a representative figure in the main text.

1~15: E8.5 embryos obtained from Nczf heterozygous intercrosses. M: DNA size marker.

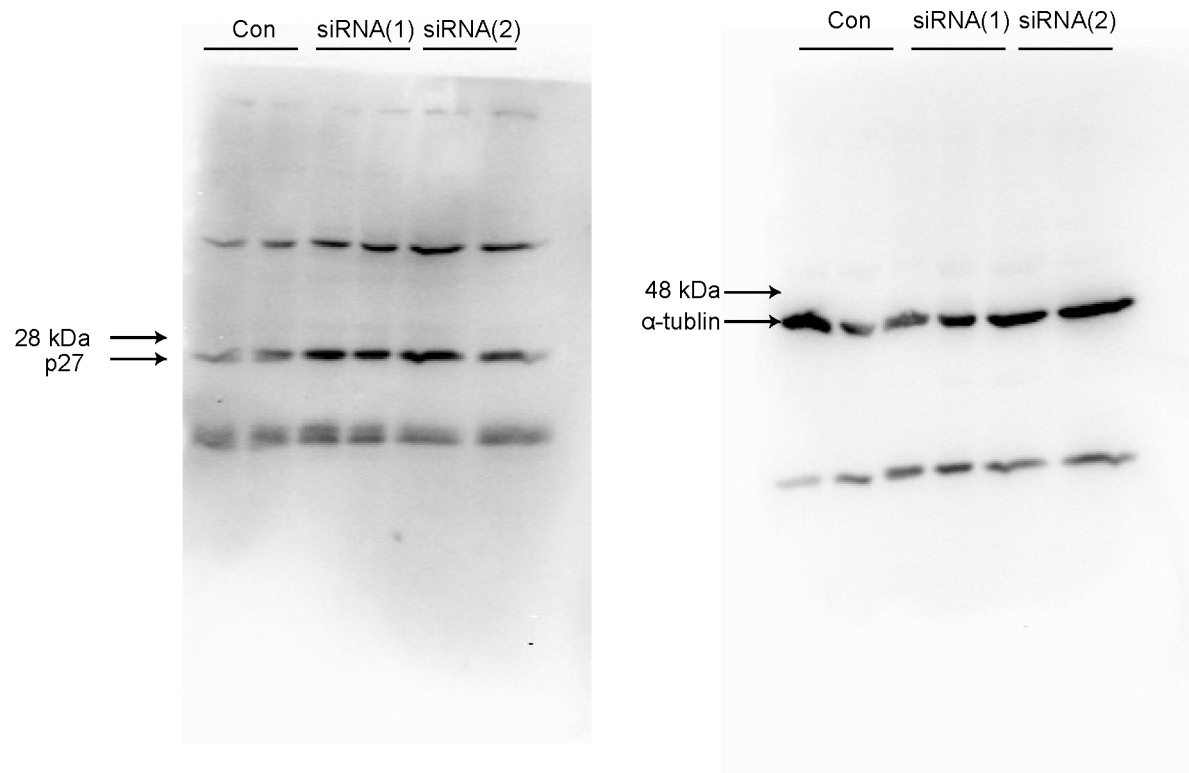


Figure S2

Whole image of western blotting. MEFs from two independent experiments in each siRNA transfection were examined by Western blot analysis. Left figure indicates p27 expression. Right figure shows tubulin expression as a loading control.